

Product datasheet for RC203500L1V

OriGene Technologies, Inc.

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IRF1 (NM_002198) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: IRF1 (NM_002198) Human Tagged ORF Clone Lentiviral Particle

Symbol: IRF1

Synonyms: IRF-1; MAR

Mammalian Cell None

Selection:

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM 002198

ORF Size: 975 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC203500).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: <u>NM 002198.1</u>

 RefSeq Size:
 3567 bp

 RefSeq ORF:
 978 bp

 Locus ID:
 3659

 UniProt ID:
 P10914

 Cytogenetics:
 5q31.1

Lytogenetics: 5q31.7

Domains: IRF

Protein Families: Druggable Genome, Transcription Factors





ORIGENE

MW: 36.5 kDa

Gene Summary:

The protein encoded by this gene is a transcriptional regulator and tumor suppressor, serving as an activator of genes involved in both innate and acquired immune responses. The encoded protein activates the transcription of genes involved in the body's response to viruses and bacteria, playing a role in cell proliferation, apoptosis, the immune response, and DNA damage response. This protein represses the transcription of several other genes. As a tumor suppressor, it both suppresses tumor cell growth and stimulates an immune response against tumor cells. Defects in this gene have been associated with gastric cancer, myelogenous leukemia, and lung cancer. [provided by RefSeq, Aug 2017]